CLAINS:

A compound of the formula

wherein A is an L, D or DL amino acid selected from the (Ala), valine (Val), group consisting of Alanine phenylalanine (Phe), para-chlpro-phenylalanine (p.Cl Phe), tryptophan (Trp), proline (Pto), serine (Ser), Threonine (Thr), tyrosine (Tyr), glutamic acid (Glu), beta alanine (Beta Ala), -aminobutyric \acid (Abu), N-methylalanine (N-Me Ala), 5-fluorotryptopham (5-F Trp), 5-bromotryptophan (5-Br Trp), 5-chlorotryptophan (5-Cl Trp), their acetylated derviatives or a pharmaceutically acceptable acid addition salt thereof;

B is an L, D or DL amino adid selected from the group consisting of threonine amid (Thr NH2), valine amide (Val NH2), proline amide (Pro NH2), hydroxyproline amide (HO Pro NH2), serine amide (Ser NH2), tyrosine amide (Tyr NH2, tryptophan amide (Trp NH2), 5-fluorotryptophan amide (5-F Trp NH2), formyl tryptophan amide (For Trp NH2), alanine amide (Ala NH2), glycine amide (Gly NH2) and methylalanine amide (Me Ala NH2);

X is L-phenylalanine (L-Phe) or L-tyrosine (L-Tyr);

Y is L-threonine (L-Thr) or L-valine (L-Val);

Z is L, D or DL 5-halo-tryptopham, in which the halogen (Halo-) is fluorine, chlorine, bromine or iodine, or Dtryptophan (D-Trp); and

C" and C' are L or D-cysteine (Cys), \sim -aminobutyric acid (Abu), aspartic acid (Asp) or lysine (Lys);

the connecting line between C" and C' signifies a bridge selected from the group consisting of carbon/carbon, carbon/sulfur, sulfur/sulfur and amide bridges; and the pharmaceutically acceptable acid addition \$alts thereof.

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A compound according to Claim 1 wherein
                C" is Cys;
                X is Phe;
                Y is Thr; and
                3. A compound according to Claim X wherein
           ि, ५ C" is Cys;
               X is Tyr;
                Z is D-Trp;
               Y is Val; and
            , ∰ C' is Cys.
                4. A compound according to Claim 1 wherein
              C" is Cys;
X is Phe;
               Y is Thr; and
B
                   A compound according to Claim {\mathscr X} wherein
           i di C" is Cys;
              X is Tyr;
               Y is Val; and
             . ⊲C' is Cys.
               6. A compound according to Claim 1 wherein
              C" is Lys;
              X is Tyr;
Z is D-Trp;
               Y is Val; and
                C' is Asp.
               7. A compound according to Claim which is
          D-Phe-Cys-Tyr-D-Trp-Lys-Val-Cys-Thr-NH2.
                8. A compound according to Claim 1 which is
           Ac-p-C1-D-Phe-Cys-Phe-D-Trp-Lys-Thr-Cys-Thr-NII2.
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A compound according to Claim which is В P.C. - Aus: 4/18/2 OSCI-D-Phe-Cys-Che-D-Trp-Lys-Cys-Cys-Mi-NH2 TOYZOX 10. A compound according to Claim which is TOULIS ACOSED-D-Phe-Cys-Tyr-D-Trp-Lys-Val-Cys-Thr-ill2. 1. A compound according to Claim & which is Ac-D-Phe-Lys-Tyr-D-Trp-Lys-Val-Asp-Thr-NH2 (amide bridge). An octapeptide (reduced form) of the formula A - C'' - X - Z - Lys - Y - C' - Bwherein A, C", X, Z, Y, C' and B are as defined in Claim -10' A compound according to Claim which is: D-Phe-Cys-Tyr-D-Trp-Lys-Val-Cys-Thr-NH2. 14. A compound according to Claim 12 which is: D-Phe-Cys-Tyr-D-Trp-Lys-Val-Cys-Ser-NH2. 12
15. A compound according to Claim 12 which is: D-Phe-Cys-Tyr-D-Trp-Lys-Val-Cys-Trp-NH2. R.C. AVS; 4-18. A process for \preparing an octapeptide of 16. A - C'' - X - Z - Lys - Y - C' - Bin which A, C", X, Z, Y, C' and B are as defined in Claim 1 which comprises oxidizing a corresponding peptide of formula (II) A - C" - X - Z - Lys - Y + in which A, C", X, Z, Y, C' and B are as defined above. 17. The process of Claim \6 wherein the peptide of formula (I') is oxidized with potassium ferricyanide, iodine, oxygen or air.

18. A pharmaceutical composition which comprises an octapeptide of Claim 1, its reduced form or a pharmaceutically acceptable acid addition salt thereof in a pharmaceutically acceptable liquid or solid carrier thereof.

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19. A pharmaceutical composition of Claim 18 which comprises an octapeptide of Claim 1, its reduced form or a pharmaceutically acceptable acid addition salt thereof, encapsulated in poly (d,1-lactide-co-glycolide) microcapsules.

20: A method of treating excess release of growth hormone, gastrointestinal disorders, cancer and diabetes in a mammal in need of such therapy which comprises administering to said mammal an effective dose of octapeptide of Claim its reduced form, or a pharmaceutically acceptable acid addition salt thereof.

(R.C.-AVP) B) 2+ A COMPOUND ACCORDING TO

(R.C.-AVS)

(R.C.-AVS)

D. Phe-Cys-Tyr-D-Typ-Lys-Val-Cys-Tyr-NH2

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